



Protocol Link

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For more information contact:

Adriane Johnson

847.549.0390, ext. 209

ajohnson@protocollink.com

Dr. Daniel Hoch to Present at Research and Lab Services Forum Hosted by Pfizer

VERNON HILLS, IL – (October 20, 2008) Protocol Link, Inc., a leading project management and consulting firm to the FDA-regulated life sciences industry, today announced that Daniel J. Hoch, Ph.D., Vice President of Project Services, has been invited to present at the inaugural Research and Lab Services Forum, which will be held on November 12, 2008, from 9:00 a.m. to 4:30 p.m. at Pfizer's PGRG Global Headquarters located in New London, Connecticut.

The Forum will attract fellow leaders in Facilities Operations and Management within the Pharmaceutical Industry and focus on sharing perspectives and discussing trends in effective management of the Laboratory space. A cohort of industry experts will provide presentations highlighting their perspectives on the topic and help moderate the discussion.

Dr. Hoch will provide expertise and present on Trends in Laboratory Services and Compliance. He will focus specifically on regulatory compliance issues and current trends within research and laboratory services, and share insight into ensuring ongoing compliance.

"It is imperative that professionals working in the industry are abreast of current trends in quality and compliance requirements that impact their day-to-day responsibilities and decision-making, as well as the importance of creating an environment of full compliance," said Dr. Hoch. "I'm honored to be a part of this Forum, which enables professionals to engage in discourse about research and laboratory services and help shape the thinking on this topic for the future."

Dr. Hoch has more than 22 years experience in the pharmaceutical and biotechnology industries. He has held significant management positions in QA/QC for major pharmaceutical companies and has expertise in process and analytical test method validation, laboratory management, and global compliance requirements.

About Protocol Link

With an established client base worldwide, Chicago-based Protocol Link, Inc., is a leading project management and consulting firm dedicated to providing comprehensive regulatory compliance, quality assurance, cGMP documentation, validation, and technology services to FDA-regulated life sciences companies worldwide. Founded in 1996, Protocol Link's mission is to provide customer-focused services with ethical conduct, mutual trust, and personnel empowerment. Additional information can be obtained by visiting Protocol Link's Web site at: www.protocollink.com.

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